

ATTORNEY'S DOCKET NO: 23999

U.S. DEPARTMENT OF COMMERCE, PATENT AND TRADEMARK OFFICE		DATE: 4 October 1999 (4.10.1999)
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		U.S. APPLICATION NO. (if known): 09/402362
INTERNATIONAL APPLICATION NO.: PCT/AU98/00234	INTERNATIONAL FILING DATE: 7 April 1998 (07.04.1998)	PRIORITY DATE CLAIMED: 7 April 1997 (07.04.1997)
TITLE OF INVENTION: FOOD GRADE WAX AND PROCESS FOR PREPARING SAME		
APPLICANT(S) FOR DO/EO/US: VALIX, Marjorie Gan		
<p>Applicant hereby submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:</p> <ol style="list-style-type: none"> 1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. 2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. 3. <input checked="" type="checkbox"/> This express request to begin national examination procedures (35 USC 371(f)) at any time rather than delay examination until the expiration of the time limit set in 35 USC 371(b) and PCT Articles 22 and 39(1). 4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. 5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2)): <ol style="list-style-type: none"> a. <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). b. <input checked="" type="checkbox"/> has been transmitted by the International Bureau. c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US) 6. <input checked="" type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)). 7. <input type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)) <ol style="list-style-type: none"> a. <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau). b. <input type="checkbox"/> have been transmitted by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input checked="" type="checkbox"/> have not been made and will not be made. 8. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). 9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). 10. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)). <p>ITEMS 11. TO 16. BELOW CONCERN OTHER DOCUMENT(S) OR INFORMATION INCLUDED:</p> <ol style="list-style-type: none"> 11. <input checked="" type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. 12. <input checked="" type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. 13. <input checked="" type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment 14. <input type="checkbox"/> A substitute specification. 15. <input type="checkbox"/> A change of power of attorney and/or address letter. 16. <input checked="" type="checkbox"/> TRANSMITTAL FORM; WIPO APPLICATION WO 98/45390 CONSISTING OF 11 PAGES SPECIFICATION, 2 PAGES WITH 17 CLAIMS AND 1 COVER SHEET CONTAINING THE ABSTRACT; PCT REQUEST FORM PCT/RO/101; PCT/ISA/210 INTERNATIONAL SEARCH REPORT; PCT/IPEA/401 DEMAND; PCT/IPEA/408 WRITTEN (FIRST); RESPONSE TO FIRST WRITTEN OPINION; PCT/IPEA/408 (SECOND); RESPONSE TO SECOND WRITTEN OPINION WITH 2 AMENDED CLAIM SHEETS FOR PAGES 12 AND 13; PCT/IPEA/409 INTERNATIONAL PRELIMINARY EXAMINATION REPORT WITH ARTICLE 34 AMENDMENTS FOR PAGES 12 AND 13 		

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17. <u>xx</u> The following fees are submitted: Basic National Fee (37 CFR 1.492(a)(1)-(5): Search Report has been prepared by the EPO or JPO:.....\$840.00 International preliminary examination fee paid to USPTO (37 CFR 1.482).....\$670.00 No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)).....\$760.00 Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO.....\$970.00 International preliminary examination fee (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4).....\$ 96.00 ENTER APPROPRIATE BASIC FEE AMOUNT =	<u>CALCULATIONS</u> \$ 970.00 \$ 970.00	<u>PTO USE ONLY</u>
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Surcharge of \$130.00 for furnishing the oath or declaration later than <u> 20 </u> <u> 30 </u> months from the earliest claimed priority date (37 CFR 1.492(e)).	\$	
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CLAIMS	NO. FILED	NO. EXTRA	RATE		
TOTAL	<u>17</u> -20=	0	X \$ 18.00	\$	0.00
INDEPENDENT	<u>2</u> - 3=	0	X \$ 78.00	\$	0.00
Multiple dependent claims(s) (if applicable)			+ \$250.00	\$	0.00
TOTAL OF ABOVE CALCULATIONS =				\$	970.00
Reduction by ½ for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28).				\$	0.00
SUBTOTAL =				\$	970.00
Processing fee of \$130.00 for furnishing the English translation later than <u> 20 </u> <u> 30 </u> months from the earliest claimed priority date (37 CFR 1.492(f)).				\$	0.00
TOTAL NATIONAL FEE =				\$	970.00
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$	40.00
TOTAL FEES ENCLOSED =				\$	1010.00
				Amount to be: refunded	\$
				charged	\$

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a. ☒ A check in the amount of \$970.00 to cover the Government Filing Fee, and a check in the amount of \$40.00 for the recordation of the Assignment, is enclosed.

b. ☐ Please charge my Deposit Account No. 14-0112 in the amount of \$_____ to cover the above fees. (A duplicate copy of this sheet is enclosed.)

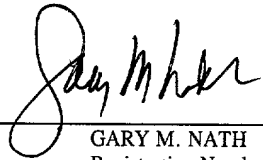
c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 14-0112.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed to request that the application be restored to pending status.

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Rev. 02/98

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

VALIX, Marjorie Gan

International Application No. PCT/AU98/00234

Serial No. NOT YET ASSIGNED

International Filing Date: 7 April 1998 (7.04.98)

Filed: October 4, 1999

For: FOOD GRADE WAX AND PROCESS FOR PREPARING SAME

TRANSMITTAL LETTERAssistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Submitted herewith for filing in the U.S. Patent and Trademark Office is the following:

- (1) Transmittal Letter
- (2) Transmittal Letter To U.S. Designated/Elected Office (DO/EO/US) Concerning Filing under 35 U.S.C. 371
- (3) WIPO Publication No: WO 98/45390
International Publication Date: 15 October 1998 (15.10.98)
which consists of 14 pages including:
 - 11 pages Textual Specification
 - 2 pages of 17 claims
 - 1 Cover Sheet Containing the Abstract
- (4) Executed Declaration
- (5) Assignment with recordation cover sheet
- (6) PCT/RO/101 Request
- (7) PCT/ISA/201 International Search Report
- (8) PCT/IPEA/401 Demand
- (9) PCT/ISA/408 Written Opinion (First)
- (10) Response to Written Opinion
- (11) PCT/ISA/408 Written Opinion (Second)
- (12) Response to Written Opinion with 2 amended claim sheets for page 12 and 13
- (13) PCT/IPEA/409 International Preliminary Examination Report with Article 34 Amendments for pages 12 and 13
- (14) Information Disclosure Statement with 5 references
- (15) PTO Form-1449
- (16) Check No. 11950 \$ 970.00 for Government Filing Fee
- (17) Check No. 11952 \$ 40.00 for the Recordation of the Assignment
- (18) Early Notification Serial Number Postcard

Date: October 4, 1999

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Respectfully submitted,

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09/402362

FOOD GRADE WAX AND PROCESS FOR PREPARING SAME
TECHNICAL FIELD

This invention relates to a wax composition suitable for use in
comestibles. The invention also relates to a process for preparing the
5 composition.

BACKGROUND ART

Many comestibles include waxes added during preparation of the
comestible. For example, wax is a component of chewing gum base. Waxes
can also be used as protective coatings on comestibles such as cheeses and
10 fruits. The waxes used for such purposes are typically mineral waxes such as
montan wax extracted from lignites, peat waxes, ceresin wax and petroleum
waxes. Among these mineral waxes, it appears that only petroleum based
waxes are used in food applications: microcrystalline waxes, high melting
point waxes and high sulfur microcrystalline waxes in particular. The US
15 Food and Drug Administration (FDA) has established regulations for the use
of petroleum wax (see 21 CFR 172.886 and 178.3710) and Japanese
authorities consider petroleum waxes to be natural products and allows their
use in products such as chewing gum. Although montan wax is not used
directly in food applications, it is used in plastic processing such as plastic
20 containers and wrappings which will come or may come into contact with
food.

Use of mineral waxes in comestibles is undesirable. Mineral waxes
are extracted from coal and crude petroleum oil. These raw materials contain
organic chemicals which are toxic to humans. The food applicability of the
25 waxes depends on the degree of refining or purification achieved and its
usage has been regulated according to the specifications provided by
authorities such as Ministry of Agriculture, Fisheries and Food, UK (The
Mineral Hydrocarbons in Food Regulation, SI 1966 No. 1073. This regulation
applies to England and Wales only, though similar regulations apply to
30 Scotland and Northern Ireland). The refining achieved has been acceptable
to food regulating authorities. However, recent studies have suggested
toxicological effects of petroleum based waxes ("Recommendations on the
use of mineral hydrocarbon in food", Food Advisory Committee 8/93, UK) and

the hydrocarbon imparted by packaging using mineral hydrocarbon waxes on food (Progress Report of the Working Party on Chemical Contaminants from Food Contact Materials: 1988 to 1992; Food Surveillance Paper No. 38, Ministry of Agriculture of Fisheries and Food). The Ministry of Agriculture, Fisheries and Food is at present reviewing studies on the toxicological impact of using petroleum based waxes as food additives and as a processing aid to update regulation of the usage of these waxes in food applications (Ministry of Agriculture of Fisheries and Food News Release 278/93 of 19 August 1993).

Waxes of plant origin are known. Indeed, the palm *Copernicia cerifera* is a source of the common wax, carnauba. Waxes can also be extracted from sugar cane and rice. Subject to the conditions used for extraction, waxes of plant origin should provide an alternative to mineral waxes for use in comestibles.

Sugar production results in a number of by-products, one of which is mill mud. Mill mud comprises crude wax and fats, fibre, sugar, crude protein and ash (SiO_2 , CaO , P_2O_5 and MgO). A crude cane wax can be extracted from mill mud. However, the crude wax is unsuitable for use in comestibles as it has a foul odour and taste and is dark green to brown in colour due to the presence of contaminants. US Patent No. 2,464,189 describes a process for the refining of sugar cane wax. However, wax produced by this process is unsuitable for use in comestibles for the following reasons:

1) The refining process is not complete. The process is only a fractionation step which removes a resinous fraction (pitch) from the sugar cane wax. The wax colour still has to be removed and stabilised. The patentees suggested further processing—for example, bleaching with acid, decolourisation, emulsification (see Example 3, line 40).

2) The reagent used in the process (acetone) is not food grade.

3) Bleaching uses reagents such as chromic and sulfuric acid which are not food grade reagents.

Since by-products of sugar production are plentiful in countries such as Australia, it would be desirable to have a process for producing a wax from such by-product (i.e., mill mud) suitable for use in comestibles.

SUMMARY OF THE INVENTION

The object of this invention is to provide a sugar cane wax composition, and a process for preparing the composition, which wax is suitable for use in comestibles.

5 In one aspect, the invention provides a wax composition comprising on a weight basis: wax esters, 6.2–11%; aldehydes, 2.8–9.5%; triglycerides, 0–3.0%; alcohols, 1.8–44.5%; and, free fatty acids, sterols and polar lipids, 36.8–87.2%.

10 In a second aspect, the invention provides a process for preparing a wax composition from crude sugar cane wax, the process comprising the steps of:

- i) heating a solution of the crude wax with a lower alcohol as solvent at the boiling point of the solvent;
- 15 ii) allowing phase separation of the solution from (i) and decanting the upper phase while hot;
- iii) allowing the separated phase from (ii) to cool and separating crystallised wax from the solvent;
- iv) repeating steps (i) to (iii) using the wax from (iii) until all pitch has been removed from the wax;
- 20 v) heating the wax to between 90 and 140°C and oxidising molten wax with oxidising material; and
- vi) continuing the heating under an inert gas on completion of the oxidation step until intermediate peroxide products are removed.

25 In other aspects, the invention provides the wax composition product of the process according to the second aspect and comestibles which include a wax composition according to the first aspect or as the product of the process according to the second aspect.

BEST MODE AND OTHER MODES FOR
CARRYING OUT THE INVENTION

30 The inventor has found that a wax composition suitable for inclusion in comestibles can be obtained from sugar cane. The composition is essentially odourless and colourless, desirable properties for compositions used as a comestible base or for coating comestibles.

Typical components of the wax composition according to the first aspect of the invention are set out in Table I below. The numbers in the table are the numbers of carbon atoms typically found in a member of a functional group. However, it will be appreciated that a member of a group may have a number of carbon atoms falling outside the indicated range. Members of groups are mostly straight chain saturated and unsaturated hydrocarbons.

Table I
Components of Groups of The Wax Composition

Functional Groups	Typical Components
Wax Esters	C16 (fatty acid)–C24(alcohol)–C16 (fatty acid)–C36 (alcohol)
Aldehydes	C28–C36
Tri-Glycerides	T48–T54 (total number of carbon in the acyl group) or C16–C18 (number of carbons in each acyl group)
Alcohols	C24–C36
Free Fatty Acid	C24–C36
Alkanes	C25–C35

The polar lipids in the wax are essentially amphipathic molecules, having a hydrophobic fatty acid part and a hydrophilic domain. The three which are commonly found are *phosphoglycerides*, in which fatty acids are esterified with an alcohol (glycerol) which contain a phosphate group, *glycosyl diglycerides* in which the fatty acids are esterified with an alcohol (glycerol) which contains a carbohydrate (sugars) and *sphingolipids* in which the fatty acids are esterified with an alcohol (glycerol) which contains amino groups.

As indicated above in the description of the second aspect, the method of refining the crude sugar cane wax involves heating the crude wax with an organic solvent to allow the pitch and the paler wax fraction to separate. These form two distinct phases, which can be separated by decantation. The paler wax fraction is cooled to allow the wax to crystallise and separate from the oil which remains soluble in the organic solvent. The wax is filtered until

dry. The wax is melted over a hot bath and oxidised by blowing fine air bubbles through a sparger. This is continued until oxidation is complete noticeable by lack of any further increase in temperature—arising from the exothermic nature of the reaction—and colour change.

5 The solvent used in step (i) of the process is typically ethanol or isopropanol which have boiling points of 78.5°C and 82.4°C, respectively. The crude wax is typically combined with solvent at a ratio of one part (by weight) wax to nine parts solvent although ratios of 1:8 to 1:20 can also be used. Heating can be for 5 to 60 minutes although a heating time of about 30
10 minutes is usually adequate.

As indicated above, the upper phase formed in step (ii) contains the wax of interest. The lower phase is a resinous fraction referred to as "pitch" which contains wax of lower quality.

In step (iii), filtration or centrifugation are advantageously used to
15 separate wax from oil-containing solvent. However, other methods known to those of skill in the art can be used.

Steps (i) to (iii) are repeated from 2 to typically no more than 5 times. The number of times the steps are repeated largely depends on the amount of pitch present in the crude wax, the rate with which the pitch settles and the
20 rate of wax crystallisation. It appears that pitch that does not settle fast enough is occluded within the wax crystals.

The heating of the wax in step (v) of the process is advantageously carried out under an oxygen-free gas. This allows better control over the oxidation of the wax which is not initiated until the composition reaches the
25 desired temperature. The oxygen-free gas is typically an inert gas such as nitrogen.

The oxidising material used in step (V) can be chromic acid, potassium permanganate, transition metals such as salts of noble metals such as platinum and palladium, pentavalent vanadium, cobalt (III), cerium (IV)
30 thallium (III), mercury (II), cupric solutions, specific enzymes, and oxygen gas (see R. Stewart, "Oxidation Mechanisms, Application to Organic Chemistry", W.A. Benjamin Inc., 1964). Preferred oxidising materials are air, oxygen, or mixtures of oxygen, nitrogen and ozone.

With regard to the final step of the process, step (vi), one of skill in the art would be able to determine when removal of the intermediate peroxide products is complete. Completion typically takes from 30 minutes to 2 hours. However, longer or shorter periods can be used depending on the degree of oxidation achieved. The inert gas used in this step is typically nitrogen.

In step (v) of the process, oxidation can be enhanced by using a catalyst. Suitable catalysts include cobalt or manganese borates and resins (A.J.C. Andersen, *Refining of Oils and Fats for Edible Purposes*, Second Revised Edition, P.N. Williams, ed., Pergamon Press, 1962), ferrous salts, and Fenton's reagent which consists of ferrous salts and H_2O_2 (Roger A Sheldon and Jay K. Kochi, *Metal Catalyzed Oxidations of Organic Compounds*, Academic Press, 1981).

The wax compositions obtained from step (vi) of the process according to the second aspect of the invention can be further decolourised, if desired, with adsorbents or by pitch inducement. Each of these methods will now be briefly described.

Use of Adsorbents

Suitable adsorbents include activated carbons, resins, activated alumina and silica. Carbons obtained from commercial sources are satisfactory and with a wax to carbon ratio of 1:3 white wax can be produced. It is also possible to manufacture carbons that are selective towards a particular colour.

The following are typical steps in the decolourisation of the wax composition with activated carbon:

- a) Wax from step (v) is heated with a lower alcohol as solvent at the boiling point of the solvent for 30 to 60 minutes with wax to activated carbon ratios of between 1:0.5 and 1:3.
- b) The molten slurry is filtered hot.
- c) The wax and solvent recovered is cooled until the wax crystallises and is separated by filtration.

Pitch Inducement

In the pitch inducement method, colour can be removed without the use of adsorbents. The method involves heat treatment and fractionation

which results in waxes of various intensity of colour from a golden yellow to cream.

Typical steps in the reduction of the colour of the wax composition with pitch inducement follow.

- 5 (a) Wax from step (v) is heated with a lower alcohol as solvent at the boiling point of the solvent for 30 to 60 minutes.
- (b) The phases of the solution from (a) are separated and the upper phase decanted while hot.
- (c) The separated upper phase from (b) is allowed to cool and the
10 crystallised wax separated from the solvent.
- (d) Wax from (c) is heat treated at 80 to 110°C in the absence of solvent for typically 15 minutes to 3 hours.
- (e) Steps (a) to (d) are repeated until the desired colour grade is achieved.

- 15 The crude sugar cane wax can be prepared by methods known to those of skill in the art. A suitable method is described, for example, in US Patent No. 2,508,002, the entire content of which is incorporated herein by cross-reference. A brief description of a suitable process follows.

Crude Wax Extraction

- 20 Sugar filter cake is steam heated and charged to a continuous reactor where it is mixed with solvent (naphtha) and held at the desired temperature and pressure. Wax is extracted into the solvent, which is subsequently separated from the filter cake. The separated wax-containing solvent is then passed through a flash drum and an evaporator to separate the crude wax
25 from the solvent. The resulting filter cake is then steam stripped to recover residual solvent.

The composition of crude sugar cane wax is typically as presented in Table II.

Table II
Composition of Crude Sugar Cane Wax

Component	Composition (% w/w)
Wax Ester	5.9–8.5
Alkyl Ketone	3.2–1.6
Tri-Glyceride	0–1.6
Alcohol	7.9–8.3
Free Fatty Acid/Sterol	5.9–7.8
Polar Lipid	73.2–76.1

Advantages of the wax composition according to the invention and the
5 process for preparing the composition are as follows:

- 1) The process produces a potential food grade vegetable wax product, which can be used as a replacement for mineral waxes in a number of food or non-food applications, including (but not limited to) chewing gum base, cheese coating, and fruit coating.
- 10 2) The process is simple and of low cost and enables economic use of wax for applications indicated in (1).
- 3) The wax product is colourless or has low colour (pale yellow) and little or no odour and taste.
- 4) The wax product has a hardness comparable to carnauba wax.
- 15 5) The wax product has good temperature stability as compared to other vegetable waxes, such as carnauba and rice wax.

Having broadly described the invention, examples of the preparation of wax composition will now be given.

Example 1

20 Multiple portions of wax composition were prepared as follows: one hundred grams of crude sugar cane wax was combined with 900 grams of ethanol in a round bottom flask. The mixture was heated in a heating mantle

to 78.5°C for 30 minutes under reflux. The solution was removed from the heating mantle and the phases of pitch and the paler wax were allowed to separate. The paler wax was decanted into another round bottom flask while the wax mixture was still in solution or only partially crystallised at 65 to 75°C.

- 5 The mixture was again heated to 78.5°C for 10 minutes. The paler wax was decanted to separate it from the pitch. The heating and separation processes were repeated about four times or until no visible pitch separated from the paler wax. The pitch was reheated with about 100 grams of ethanol to recover additional paler wax. This was combined with the previously collected paler wax.
- 10 The paler wax was then cooled slowly to room temperature and then in an ice bath. The cooled wax was filtered and remelted in a hot bath, in the presence of nitrogen, at a temperature between 90 and 140°C. Air or ozone was blown into the molten wax by means of a sparger until the temperature of the wax stabilised to a constant value or until there was no further visible
- 15 change in the colour of the wax. The overall process was executed within several hours depending on the temperature and air distribution.

Wax composition produced by the above process had the following properties: pale yellow in colour with a sweet smell and little or no taste. The compositions of waxes are summarised in the following table.

20

Table III

Components of Wax Compositions of Example 1

Component	Composition (% w/w)	
	Ethanol fractionated then oxidised with air	Ethanol fractionated then oxidised with ozone
Wax Ester	6.2–11	6.2–7.7
Aldehyde	8.1–9.5	2.5–9.5
Tri-Glyceride	0.5–3.0	0.5–3
Alcohols	11.5–44.5	1.8–44.5
Free Fatty Acid + Sterol + Polar Lipid	36.8–70	36.8–87.2

The make-up of the composition obtained following the oxidation step depends on the extent of oxidation. If complete oxidation is achieved, the only substantial component left will be free fatty acid. However, oxidation needs to be only to the extent that sufficient colour is removed.

5

Example 2

Multiple portions of wax composition were again prepared. One hundred grams of crude sugar cane wax was combined with 900 grams of isopropanol in a round bottom flask. The mixture was heated in a heating mantle to 82.4°C. The mixture was cooled slowly to room temperature and then in an ice bath. The wax was filtered and transferred into another round bottom flask. To the wax, 450 grams of ethanol was added and the mixture heated to 78.5°C for 30 minutes. Fractionation and oxidation of this wax was carried out as in Example 1.

Wax composition produced by the above process had the following properties: pale yellow in colour with a sweet smell and little or no taste. Compositions obtained are summarised in the following table.

15

Table IV
Components of Wax Compositions of Example 2

Component	Composition (% w/w)
Wax Ester	10.5–11
Aldehyde	7.4–8.1
Tri-Glyceride	0–0.6
Alcohols	11.5–18.8
Free Fatty Acid + Sterol + Polar Lipid	63.4–70

20

Yields of fractions produced using processes such as described in Examples 1 and 2 are presented in Table V.

Table V
Fractionation Yield

Fraction	Yield (%)	
	Example 1	Example 2
Oil	24-30	53-55
Pitch	26-35	20-25
Wax Composition	34-48	20-25

5 It will be appreciated that many changes can be made to the processes and compositions as exemplified above without departing from the broad ambit and scope of the invention.

CLAIMS

1. - A food grade wax composition comprising on a weight basis: wax esters, 6.2–11%; aldehydes, 2.8–9.5%; tri-glycerides, 0–3%; alcohols, 1.8–44.5%; and, free fatty acids, sterols and polar lipids, 36.8–87.2%.
- 5 2. A process for preparing a wax composition from crude sugar cane wax, the process comprising the steps of:
- i) heating a solution of the crude wax with a lower alcohol as solvent at the boiling point of the solvent;
 - ii) allowing phase separation of the solution from (i) and decanting
 - 10 the upper phase while hot;
 - iii) allowing the separated phase from (ii) to cool and separating crystallised wax from the solvent;
 - iv) repeating steps (i) to (iii) using the wax from (iii) until all pitch has been removed from the wax;
 - 15 v) heating the wax to between 90 and 140°C and oxidising molten wax with oxidising material; and
 - vi) continuing the heating under an inert gas on completion of the oxidation step until intermediate peroxide products are removed.
3. The process according to claim 2, wherein said lower alcohol is
- 20 ethanol or iso-propanol.
4. The process according to claim 2, wherein said crude wax is combined with solvent at a ratio of 1:8 to 1:20 by weight.
5. The process according to claim 4, wherein said ratio is 1:9.
6. The process according to claim 2 wherein in step (i) said solution is
- 25 heated for 5 to 60 minutes.
7. The process according to claim 6, wherein said solution is heated for about 30 minutes.
8. The process according to claim 2, wherein in step (iii) said separation is by filtration or centrifugation.
- 30 9. The process according to claim 2, wherein steps (i) to (iii) are repeated from 2 to 5 times.
10. The process according to claim 2, wherein in step (v) said heating is carried out under an oxygen-free gas.

11. The process according to claim 10, wherein said gas is nitrogen.
12. The process according to claim 2, wherein said oxidising material of step (v) is selected from the group consisting of air, oxygen, and mixtures of oxygen, nitrogen and ozone.
- 5 13. The process according to claim 2, wherein in step (v) said oxidation is carried out in the presence of a catalyst.
14. The process according to claim 10, wherein said catalyst is selected from the group consisting of a borate or resinate of cobalt or manganese, ferrous salts, and Fenton's reagent.
- 10 15. The process according to claim 2 comprising the further steps of:
- vii) heating wax from step (vi) with a lower alcohol as solvent at the boiling point of the solvent with activated carbon present at a wax to carbon ratio of 1:0.5 to 1:3;
 - viii) filtering the molten slurry while hot;
 - 15 ix) allowing the recovered wax/solvent mixture to cool and separating crystallised wax therefrom.
16. The process according to claim 2 comprising the further steps of:
- (vii) heating wax composition from step (vi) with a lower alcohol as solvent at the boiling point of said solvent for 30 to 60 minutes;
 - 20 (viii) allowing phase separation of the solution from (vi) and decanting the upper phase while hot;
 - (ix) allowing the separated upper phase from (viii) to cool and separating crystallised wax from said solvent;
 - (x) heating wax from (ix) in the absence of solvent for 15 minutes to 25 3 hours; and
 - (xi) repeating steps (vii) to (x) until the desired degree of decolourisation is achieved.
17. A comestible which includes the food grade wax composition of claim 1.

DECLARATION FOR PATENT APPLICATION

ATTY DOCKET
23999

As a below-named inventor, I hereby declare that:

My residence, post office address and citizenship is as stated below next to my name.

I believe I am the original, first and sole inventor of the subject matter which is claimed, and for which a patent is sought on the invention entitled:

FOOD GRADE WAX AND PROCESS FOR PREPARING SAME

the specification of which: (check one)

☒ is attached hereto.

☐ was filed on _____, as Serial No. _____
and was amended on _____ 19_____(if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the patentability of this application as defined by 37 CFR § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. § 119 of any foreign application(s) for patent or inventor's certificate listed below, and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Applications:

PO 6050
(Application No.)

Australia
(Country)

07/04/97
(Day/Month/Year Filed) Priority Claimed
☒ ☐
Yes No

(Application No.)

(Country)

/ /
(Day/Month/Year Filed) Priority Claimed
☐ ☐
Yes No

(Application No.)

(Country)

/ /
(Day/Month/Year Filed) Priority Claimed
☐ ☐
Yes No

654007 23999-0460

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I hereby appoint Gary M. Nath, Reg. No. 26,965; Irvin A. Lavine, Reg. No. 16,838; Karen Lee Orzechowski, Reg. No. 31,621; Harold L. Novick, Reg. No. 26,011; Suet M. Chong, Reg. No. 38,104; Todd . Juneau, Reg. No. 40,699; Patricia M. Drost, Reg. No. 29,790; Leigh A. Penfield, Reg. No. 37,050; Donald L. Sandler, Reg. No. 19,237; and Robert G. Lev, Reg. No. 30,280 as my attorneys to prosecute this application and transact all business in the U.S. Patent and Trademark Office connected therewith.

Direct Telephone Calls to:

Send Correspondence to:

Gary M. Nath
(202) 775-8383

Nath & Associates
6th Floor
1030 15th Street, N.W.
Washington, D.C. 20005 U.S.A.

I hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by 35 U.S.C. § 112, first paragraph, I acknowledge the duty to disclose material information as defined in 37 CFR § 1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

(U.S. Application Serial No.) (U.S. Filing Date) (Status-patented, pending, abandoned)

(U.S. Application Serial No.) (U.S. Filing Date) (Status-patented, pending, abandoned)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. § 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of sole inventor: Marjorie Gan Valix

Inventor's Signature:

Marjorie Gan Valix

Date

23/9/99

Residence: West Ryde, New South Wales, Australia

Aux

Citizenship: AustralianPost Office Address: 26 Andrews Street, West Ryde, NSW 2114, Australia

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